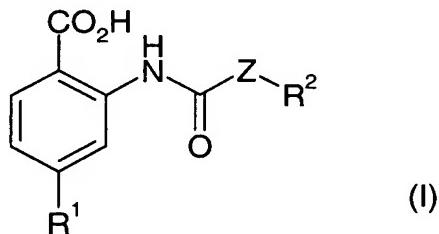


**Amendments to the claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of claims:**

1. (Currently amended): A[[n]] compound selected from: a compound of Formula (I) :



and or a salt, solvate or physiologically functional derivative thereof, wherein:

R<sup>1</sup> represents is hydrogen, halogen or C<sub>1</sub>-C<sub>3</sub>alkyl;

R<sup>2</sup> represents is a 5 or 6-member aryl, heteroaryl, heterocyclic or alicyclic ring;

Z represents is -(CH<sub>2</sub>)<sub>q</sub>- [[;]] , -CH=CH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)NH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)O- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>SO<sub>2</sub>NR<sup>3</sup>- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NR<sup>3</sup>SO<sub>2</sub>- [[;]] , -(CH<sub>2</sub>)<sub>n</sub>O- [[;]] , -C(R<sup>4</sup>R<sup>5</sup>)O- or -Y-W-X- ;

W represents is a 5 or 6-member aryl, heteroaryl, heterocyclic or alicyclic ring;

X and Y[[,]] which may are independently be present or absent, where present independently represent is -(CH<sub>2</sub>)<sub>q</sub>- [[;]] , -CH=CH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)O- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)NH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>SO<sub>2</sub>NR<sup>3</sup>- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NR<sup>3</sup>SO<sub>2</sub>- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>C(O)- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>O- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>S- or -(CH<sub>2</sub>)<sub>p</sub>O-CH<sub>2</sub>- ;

n represents an integer selected from is 2, 3 and or 4;

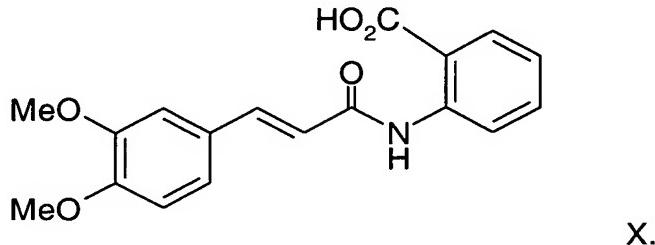
p represents an integer selected from is 0, 1 and or 2;

q represents an integer selected from is 1, 2, 3 and or 4;

R<sup>3</sup> represents is hydrogen or methyl; and

R<sup>4</sup> and R<sup>5</sup> ,which may be the same or different, are independently represent C<sub>1</sub>-C<sub>3</sub>alkyl; provided

- (i) that when R<sup>1</sup> is hydrogen, Z is -(CH<sub>2</sub>)<sub>n</sub>-, and n is 2, then R<sup>2</sup> is other than para-chlorophenyl or para-methylphenyl; and
- (ii) that a compound of Formula (I) is other than 2-(2-((4-(phenyl)phenyl)amino)acetyl)benzoic acid, 2-(2-((4-phenyl)phenoxy)acetyl)amino) benzoic acid, 2-[[(4-cyclohexylphenoxy)acetyl]amino]benzoic acid, 2-[[3-[3-(4-chlorophenyl)-1,2,4-oxadiazol-5-yl]-1-oxopropyl]amino]benzoic acid or compound X

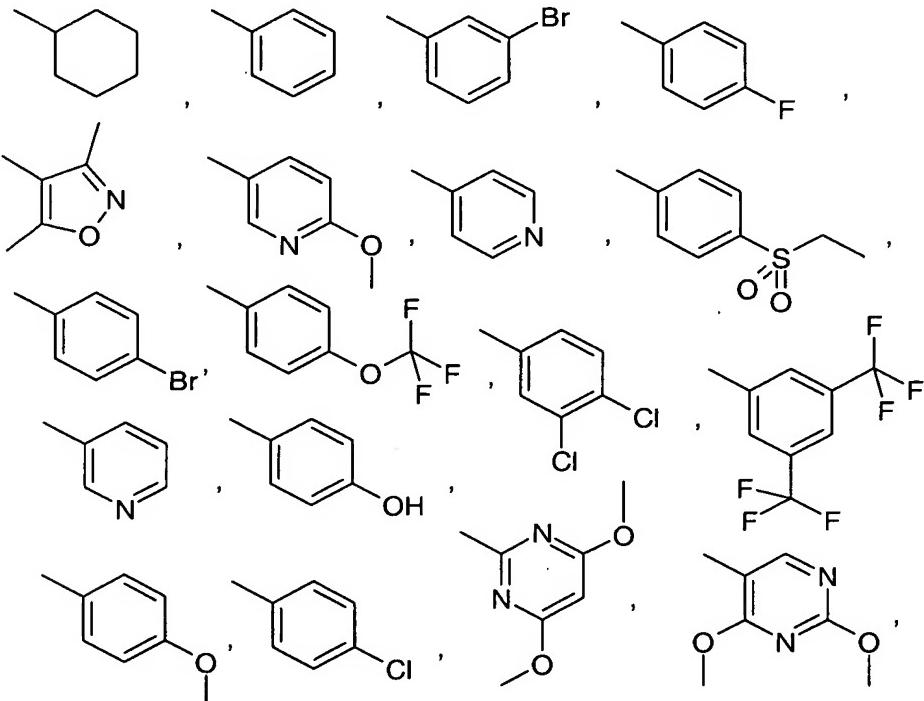


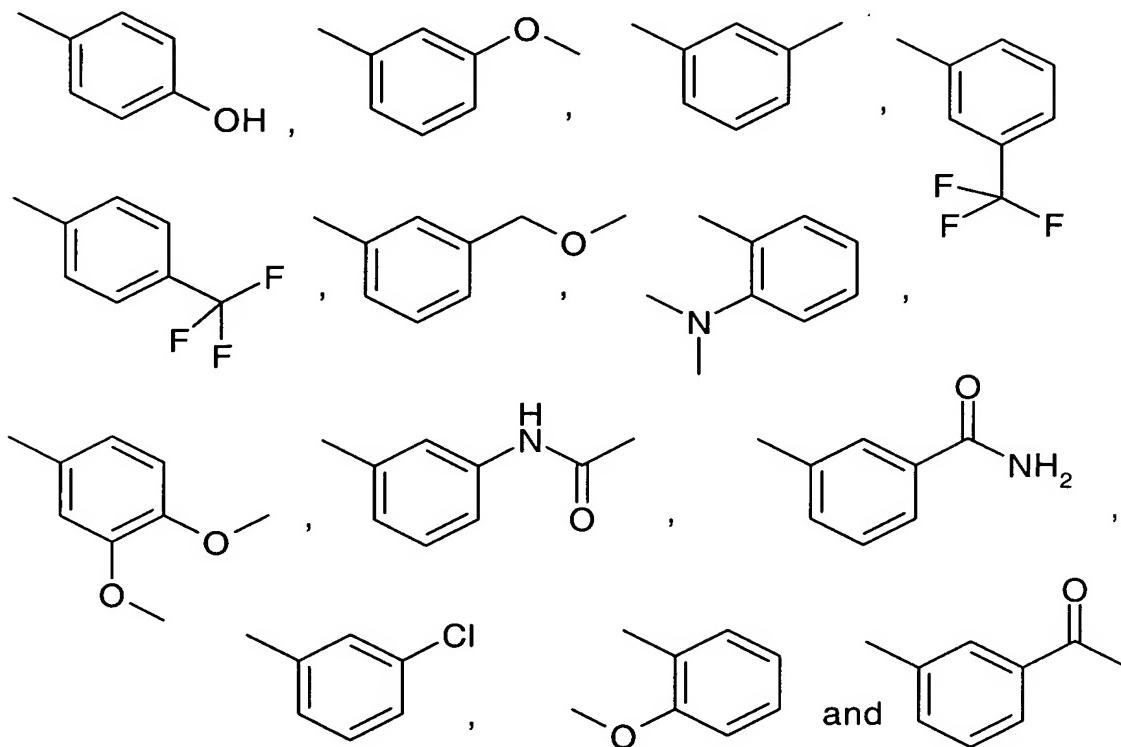
2. (Original): A compound according to claim 1 wherein R<sup>1</sup> is hydrogen or methyl.

3. (Original): A compound according to claim 2 wherein R<sup>1</sup> is hydrogen.

4. (Currently amended): A compound according to claim 1 ~~any preceding claim~~ wherein R<sup>2</sup> is cyclohexyl, phenyl, pyridinyl, pyrimidinyl, pyridazinyl and/or isoxazolyl.

5. (Currently amended): A compound according to claim 1 ~~any one of claims 1-3~~ wherein R<sup>2</sup> is selected from the group consisting of:





6. (Currently amended): A compound according to claim 1 ~~any one of claims 1-3~~ wherein R<sup>2</sup> is substituted phenyl.

7. (Currently Amended): A compound according to claim 6 wherein R<sup>2</sup> is phenyl substituted with one or two substituents ~~selected from which are~~ halogen C<sub>1-3</sub>alkyl, C<sub>1-3</sub>haloalkyl C<sub>1-3</sub>alkoxy and ~~or~~ C<sub>1-3</sub>haloalkoxy.

8. (Currently amended): A compound according to claim 1 ~~any preceding claim~~ wherein Y is -O-, -CH<sub>2</sub>- or -CH<sub>2</sub>O-.

9. (Currently amended): A compound according to claim 1 ~~any preceding claim~~ wherein X is absent or is -SO<sub>2</sub>NR<sup>3</sup>-, -NHC(O)- or -NHC(O)NH-.

10. (Currently amended): A compound according to claim 1 ~~any preceding claim~~ wherein Y is -CH<sub>2</sub>- and X is -SO<sub>2</sub>NR<sup>3</sup>-.

11. (Currently amended): A compound according to claim 1 ~~any one of claims 1-7~~ wherein Y is -O- and X is absent.

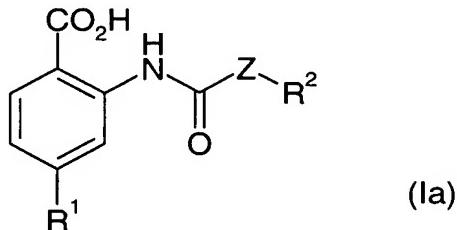
12. (Currently amended): A compound according to claim 1 ~~any preceding claim~~ wherein W is a 5 or 6 member aryl or heteroaryl ring.

13. (Original): A compound according to claim 12 wherein W is phenyl.

14. (Original): A compound according to claim 12 wherein W is a 5 member heteroaryl ring.

Claims 15-20 (Cancelled).

21. (Currently Amended): A method for the treatment of a human or animal subject having ~~disease~~ a condition characterised by under-activation of the HM74A receptor or in which activation of the receptor will be beneficial, which method comprises administering to said human or animal subject an effective amount of a compound ~~selected from:~~ a compound of Formula (Ia) :



And or a salt, solvate or physiologically functional derivative thereof, wherein:

R<sup>1</sup> ~~represents~~ is hydrogen, halogen or C<sub>1</sub>-C<sub>3</sub>alkyl;

R<sup>2</sup> ~~represents~~ is a 5 or 6-member aryl, heteroaryl, or heterocyclic or alicyclic ring;

Z ~~represents~~ is -(CH<sub>2</sub>)<sub>n</sub>- [[;]] , -CH=CH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)NH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)O- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>SO<sub>2</sub>NR<sup>3</sup>- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NR<sup>3</sup>SO<sub>2</sub>- [[;]] , -(CH<sub>2</sub>)<sub>q</sub>O- [[;]] , -C(R<sup>4</sup>R<sup>5</sup>)O- or -Y-W-X- ;

W ~~represents~~ is a 5 or 6-member aryl, heteroaryl, heterocyclic or alicyclic ring;

X and Y[[,]] ~~which may are~~ independently present or absent, where present independently ~~represent~~ is -(CH<sub>2</sub>)<sub>q</sub>- [[;]] , -CH=CH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)O- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NHC(O)NH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>SO<sub>2</sub>NR<sup>3</sup>- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NR<sup>3</sup>SO<sub>2</sub>- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>C(O)- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>NH- [[;]] , -(CH<sub>2</sub>)<sub>p</sub>O- or -(CH<sub>2</sub>)<sub>p</sub>O-CH<sub>2</sub>- ;

n ~~represents an integer selected from~~ is 2, 3 and or 4;

p ~~represents an integer selected from~~ is 0, 1 or 2;

q ~~represents an integer selected from~~ is 1, 2, 3 and or 4;

R<sup>3</sup> ~~represents~~ is hydrogen or methyl; and

~~R<sup>4</sup> and R<sup>5</sup> [[,] which may be the same or different, are independently represent C<sub>1</sub>-C<sub>3</sub>alkyl.~~

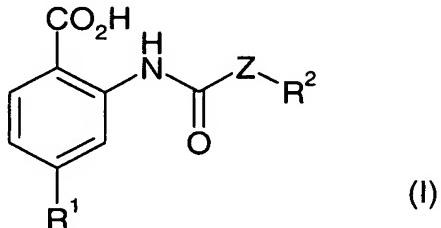
22. (Currently Amended): A method according to claim 21 wherein the condition is a disorder of lipid metabolism ~~including dislipidaemia or hyperlipoproteinæmia~~ or an inflammatory disease or condition.

23. (Currently amended): A pharmaceutical formulation comprising a compound according to claim 1 ~~any one of claims 1-14~~ in admixture with one or more physiologically acceptable diluents, excipients or carriers.

24. (Currently amended): A combination for administration together or separately, sequentially or simultaneously in separate or combined pharmaceutical formulations, said combination comprising a compound according to claim 1 ~~any one of claims 1-14~~ together with another therapeutically active agent.

25. (Currently amended): A pharmaceutical formulation comprising a compound according to claim 1 ~~any one of claims 1-14~~, plus a further active ingredient selected from the group consisting of statins, fibrates, bile-acid binding resins and nicotinic acid and one or more physiologically acceptable diluents, excipients or carriers.

26. (Currently Amended): A ~~method~~ process for the preparation of a compound of Formula (I):



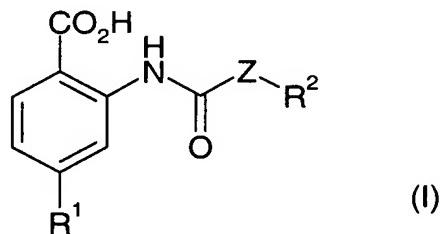
in which R<sup>1</sup> represents ~~is~~ hydrogen, Z represents ~~is~~ -Y-W-X-, Y represents ~~is~~ -(CH<sub>2</sub>)<sub>p</sub>O-, p represents ~~the integer is~~ 1, and W, X and R<sup>2</sup> are as defined in claim 1, the ~~method~~ process comprising the steps of:

- (i) amide bond formation by acetylation of an ester of anthranilic acid;
- (ii) addition of W or W-X-R<sup>2</sup> by substitution of a leaving group;
- (iii) deprotection of the anthranilic acid group;

and where desired or necessary converting a resultant free acid or base base or salt compound of Formula (I) into a physiologically acceptable salt form or free base vice versa or converting one salt form into another physiologically acceptable salt form.

27. (Currently Amended): A method process according to claim 26 where in step (ii) comprises addition of W and a further step (ii)(a), addition of R<sup>2</sup>, is included in the form of a further substitution reaction.

28. (Currently Amended): A method process for the preparation of a compound of Formula (I):



in which R<sup>1</sup>, R<sup>2</sup> and Z are as defined in claim 1, the method process comprising the steps of:

- (i) formation of an amide between the amine group of 2-amino-bezoic acid and an activated acyl transfer reagent derived from a carboxylic acid; and
- (ii) where desired or necessary converting a resultant free base or acid acid or base compound of Formula (I) into a physiologically acceptable salt form or free base vice versa or converting one salt form into another physiologically acceptable salt form.

29. (New): A method according to claim 22 wherein the disorder of lipid metabolism is dislipidaemia or hyperlipoproteinaemia.